Exhibit B



Tension-free Support for Incontinence

GYNECARE TVT™

Tension-free Vaginal Tape

GYNECARE TYT" Single Use Device GYNECARE TYT" Reusable Introducer GYNECARE TYT" Reusable Rigid Catheter Guide

GYNECARE TVT™ anordning til engangsbrug GYNECARE TVT™ indfører til flergangsbrug GYNECARE TVT™ stift guiding kateter til flergangsbrug

GYNECARE TVT** hulpmiddel voor eenmalig gebruik GYNECARE TVT** herbruikbare introducer GYNECARE TVT** herbruikbare starre kathetervoerder GYNECARE TVT** -laite, kertakäyttöinen GYNECARE TVT** -sisääriviejä, kestokäyttöinen GYNECARE TVT** -katetrinohjain, kestokäyttöinen, jäykkä

Dispositif GYNECARE TVT™ à usage unique Introducteur réutilisable GYNECARETVT™ Guide de sonde rigide réutilisable GYNECARE TVT™

GYNECARE TVT™ Einmal-Implantat GYNECARE TVT™ wiederverwendbares Einführungsinstrument GYNECARE TVT™ wiederverwendbare starre Katheterführung

Συσκευή μιας χρήσης GYNECARE TVT™ Επαναχρήσιμοποιήσιμος είσαγωγέας GYNECARE TVT™ Επαναχρησιμοποιήσιμος ακαμπτος οδηγός καθετήρα GYNECARE TVT™

Dispositivo monouso GYNECARE TY™ Introduttore riutilizzabile GYNECARE TY™ Guida rigida riutilizzabile per catetere GYNECARE TYT™

Dispositivo de utilização única GYNECARE TVT™ Introdutor reutilizável GYNECARE TVT™ Guia rígido de cateter reutilizável GYNECARE TVT™

Sistema para un solo uso GYNECARE TVT™ Introductor reutilizable GYNECARE TVT™ Guía de catéter rígida reutilizable GYNECARE TVT™

GYNECARE TVT™ äteranvändbar införare GYNECARE TVT™ återanvändbar införare GYNECARE TVT™ återanvändbar stel kateterguide



Manufactured for: ETHICON Women's Health & Urology A division of ETHICON, INC. a Johnson & Johnson company Somerville, New Jersey 08876-0151

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GYNECARE TVT™ Single Use Device GYNECARE TVT™ Reusable Introducer GYNECARE TVT™ Reusable Rigid Catheter Guide

Please read all information carefully.
Failure to properly follow instructions may result in improper functioning of the device and lead to injury.

Important:

Important:
This package insert is designed to provide instructions for use of the Tension-free Vaginal Tape single use device, Reusable Introducer, and Reusable Rigid Catheter Guide. It is not a comprehensive reference to surgical technique for correcting Stress Urinary Incontinence (SU). The device should be used only by physicians trained in the surgical treatment of Stress Urinary Incontinence and specifically in implanting the GYNECARE TVT™ Device. These instructions are recommended for general use of the device. Variations in use may occur in specific procedures due to individual technique and patient anatomy.

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DESCRIPTION (System)

GYNECARE TVT consists of the following:

GYNECARE TVT™ Single Use Device, provided sterile
(available separately)

GYNECARE TVT™ Reusable Introducer, provided non-sterile
(available separately)

GYNECARE TVT™ Reusable Rigid Catheter Guide, provided
non-sterile (available separately)

GYNECARE TVT DEVICE
The GYNECARE TVT Device is a sterile single use device, consisting
of one piece of undyed or blue (Phthalocyanine blue, color index.
Number 74160) PROLENE™ Polypropylene Mesh (tape) approximately
1/2 x 18 inches (1.1 x 45 cm), covered by a plastic sheath cut and
overlapping in the middle, and held between two stainless steel
needles bonded to the mesh and sheath with plastic collars.

PROLENE Whesh is constructed of knitted plastic collars.

PROLENE Whesh is constructed of knitted filaments of extruded polypropylene strands identical in composition to that used in PROLENE* Polypropylene Nonabsorbable Surgical Sutures. The mesh is approximately 0.027 inches (0.7 mm) thick. This material, when used as a suture, has been reported to be non-reactive and to retain its strength indefinitely in clinical use. PROLENE Mesh is knitted by a process which interlinks each fiber junction.

process which interlinks each fiber junction.

GYNECARE TVT INTRODUCER

The GYNECARE TVT Introducer is provided non-sterile and is reusable. The Introducer is made of stainless steel. It consists of two parts, a handle and an inserted threaded metal shaft. The Introducer is intended to facilitate the passage of the GYNECARE TVT Device from the vagina to the abdominal skin. It is connected and fixed to the needle, via the threaded end of the shaft, prior to inserting the needle with the tape.

GYNECARE TVT RIGID CATHETER GUIDE
The GYNECARE TVT Rigid Catheter Guide is a non-sterile reusable instrument intended to facilitate the identification of the uretria and the bladder neck during the surgical procedure. It is inserted into a Foley catheter (recommended size 18 French) positioned in the bladder via the urethra. To facilitate insertion, it can be lubricated with gel.

INDICATIONS

The GYNECARETVT Device is intended to be used as a pubourethral sling for treatment of Stress Urinary Incontinence (SUI), for female urinary incontinence resulting from urethral hypermobility and/or intrinsic sphincter deficiency. The GYNECARETVT Introducer and Rigid Catheter Guide are available separately and are intended to facilitate the placement of the GYNECARETVT Device.

- INSTRUCTIONS FOR USE

 1. The procedure can be carried out under local anesthesia, but it can also be performed using regional or general anesthesia.

 2. Before the patient is prepped and draped, she should be placed in the lithotomy position taking care to avoid hip flexion greater than 60°.

 3. Insert an 18 French Folay catheter and leave that a card.

- The procedure can be carried out under local anesthesia, but it can also be performed using regional or general anesthesia.
 Before the patient is prepped and draped, she should be placed in the lithotomy position taking care to avoid hip flexion greater than 60°.
 Insert an 18 French Foley catheter and leave it to open drainage.
 At the level of the mid urethra, inject a small amount of local anesthesia submucosally to create a space between the vaginal wall and the periverthal fascia. The extent of dissection required for placement is minimal. Only a small paraurethal incision is required over the mid urethra to position the tip of the Needle. Using forceps, grasp the vaginal wall at each side of the urethra. Using a small scalpel, make a sagittal incision about 1.5 cm long starting approximately 1.0 cm cephalad from the urethral meatus. This incision will be positioned over the mid-urethral 20ne and will allow for subsequent passage of the Implant.
 With a small pair of blunt scisors, make two small paraurethral lateral dissections (approximately 0.5 to 1.0 cm) to accommodate the tips of the Needle.
 Identify the two Needle exit sites, which should be 2-2.5 cm on each side of the midline, immediately above the pubic symphysis. Either mark these sites or, if desired, place two small 3-4 mm transverse stab incisions at the intended exit site. In order to avoid the inferior epigastric vessels it is important that the exit sites is both to avoid the exit sites for the Needle passages be near the midline and close to the superior aspect of the public bone, in order to avoid anatomic structures in the abdomen, inguinal area and lateral pelvic sidewall. If retropubic inflitration of local anesthesia is not performed then consider inflitrating the retropubic space with two injections of normal salien on either side of midline. It is important that the exit sites pass an 18 gauge needle along the back of the pubic bone until the tip of the needl

- At this point, the non-dominant hand is moved from the vagina to the suprapubic exit site. The Needle tip is guided through the retropubic space staying as close to the back of the pubic symphysis as possible. This is achieved by lowering the GYNECARE TVI Introducer Handle, thereby pressing the Needle tip against the back of the pubic Done.
 During passage through the retropubic space aim the Needle tip towards the pre-marked abdominal exit site.
 When the needle tip has reached the abdominal incision unscrew the GYNECARE TVI Introducer from the Needle. Before the Implant is pulled into place, remove the 18 French Foley catheter and perform a cystoscopy (70 degree lens recommended).
 Once bladder integrity is confirmed, pull the Needle upward to bring the Implant out through the abdominal exit site. Clamp the Implant just below the Needle. Cut the Implant between the connection to the Needle and the clamp.
 The procedure is now repeated on the patient's other side while repeating steps 9 15. NOTE: IN ORDER TO MINIMIZE THE RISK. OF BLADDER INJURY, IT IS IMPORTANT THAT THE BLADDER BE DISPLACED TOTHE CONTRALATERAL SIDE USING THE MANEUVERS OUTLINED IN STEP 10.
 The ends of the implant are then pulled upward to bring the implant (slingl) loosely, i.e., without tension, under the midurethra. Adjust the Implant so that leakage is reduced, allowing only a few drops of urinary leakage to occur under stress. For this, use patient feedback, i.e. coughing with a full bladder (approximately 300 ml.).
 Grasp the Implant the sharts that surround the Implant with clamps, taking care not to grasp the Implant. Then individually remove the Implant Sheaths by gently pulling up on the clamps, away from the abdomen, one at a time. To avoid putting tension on the Implant Sheaths by gently pulling up on the clamps, survey from the abdomen one at a time. To avoid putting tension on the Implant Sheaths.
 NOTE: Premature removal of the shea

CONTRAINDICATIONS

As with any suspension surgery, this procedure should not be performed in pregnant patients. Additionally, because the PROLENE Mesh will not stretch significantly, it should not be performed in patients with future growth potential including women with plans for future pregnancy.

- patients with future growth potential including women with plans for future pregnancy.

 WARNINGS AND PRECAUTIONS

 Do not use GYNECARE TVT procedure for patients who are on anticoagulation therapy.

 Do not use GYNECARE TVT procedure for patients who have a urinary tract infection.

 Users should be familiar with surgical technique for bladder neck suspensions and should be adequately trained in implanting the GYNECARE TVT System before employing the GYNECARETVT Device. It is important that the tape be located without tension under mid-urethra.

 Acceptable surgical practice should be followed for the GYNECARETVT procedure as well as for the management of contaminated or infected wounds.

 The GYNECARETVT procedure should be performed with care to avoid large vessels, nerves, bladder and bowel. Attention to local anatomy and proper passage of needles will minimize risks.

 Retropubic bleeding may occur post-operatively. Observe for any symptoms or signs before releasing the patient from the hospital.

 Gystoscopy should be performed to confirm bladder integrity or recognize a bladder perforation.

 The Rigid Catheter Guide should be gently pushed into the Foley catheters or that the catheter guide does not extend into the holes of the Foley catheter.

 When removing the Rigid Catheter Guide, open the handle completely so that the catheter remains properly in place.

- Do not remove the plastic sheath until the tape has been properly postioned.
 Ensure that the tape is placed with minimal tension under mid-urethra.
 PROLENE Mesh in contaminated areas should be used with the understanding that subsequent infection may require removal of the material.

 The patient should be counseled that future pregnancies may negate the effects of the surgical procedure and the patient may again become incontinent.

 Since no clinical experience is available with vaginal delivery following the GYNECARE TVT procedure, in case of pregnancy delivery via cesarean section is recommended to refrain from heavy lifting and/or exercise (i.e., cycling, jogging) for at least three to four weeks and intercourse for one month. The patient can return to other normal activity after one or two weeks.
 Should dysuria, bleeding or other problems occur, the patient is instructed to contact the surgeon immediately.

 All surgical instruments are subject to wear and damage under normal use. Before use, the instrument should be visually inspected. Defective instruments or instruments that appear to be corroded should not be used and should be discarded.

 As with other incontinence procedures, de novo detrusor instability may occur following the GYNECARE TVT procedure. To minimize this risk, make sure to place the tape tension-free in the mid-urethral position.

 Do not contact the PROLENE Mesh with any staples, clips or clamps, as mechanical damage to the mesh may occur.

 Do not resterilize GYNECARE TVT Device. Discard opened, unused devices.

 ADVERSE REACTIONS

 Punctures or accertations of vessels, nerves, bladder or bowel may

- unused devices.

 ADVERSE REACTIONS

 Punctures or lacerations of vessels, nerves, bladder or bowel may occur during needle passage and may require surgical repair.

 Transitory local irritation at the wound site and a transitory foreign body response may occur. This response could result in extrusion, erosion, fistula formation and inflammation.

 As with all foreign bodies, PROLENE Mesh may potentiate an existing infection. The plastic sheath initially covering the PROLENE Mesh is designed to minimize the risk of contamination.

 Over correction, i.e., too much tension applied to the tape may cause temporary or permanent lower urinary tract obstruction.

 ACTIONS

ACTIONS

Animal studies show that implantation of PROLENE Mesh elicits a minimal inflammatory reaction in tissues and stimulates the deposition of a thin fibrous layer of tissue that can grow through the interstices of the mesh, thus incorporating the mesh into adjacent tissue. The material is not absorbed, nor is it subject to degradation or weakening by the action of tissue enzymes.

INTRICATIONS FOR CLEANING PERSARIE INSTRUMENTS

by the action of tissue enzymes.

INSTRUCTIONS FOR CLEANING REUSABLE INSTRUMENTS

(GYNECARE TVT Introducer, GYNECARE TVT Rigid Catheter

Guide)

To ensure the reliability and functionality of the GYNECARE

TVT Introducer and GYNECARE TVT Rigid Catheter Guide, clean

the instruments before initial use and after each procedure.

The following are suggested manual and automated methods

for cleaning the instruments. Prior to cleaning, the GYNECARE

TVT Introducer should be separated into its component parts

(handle and threaded shaft). The Introducer is reassembled after

cleaning and before sterilization.

Manual Method:

- Manual Method:

 1. Soak the instrument components in an enzyme cleaner suitable for stainless steel instruments.

 2. Wash in a surgical destregent and disinfecting solution at a temperature of 86°F to 95°F (30°C to 35°C). Remove any contamination from body fluids or tissues using a soft brush.

 3. Place the instrument components in an ultrasonic bath with fresh detergent solution for approximately 10 minutes or follow the instructions below if using an automatic washing cycle.

 4. Rinse thoroughly in a stream of fresh tap water followed by towel drying. The instrument components may be treated with instrument lubricant.

Automated Method:
Automatic washing cycles are suitable for stainless steel instruments.
One recommended cycle is described below:
- Rinse-Met Cycle Cold Water - 1 minute
- Wash 176° (80°C) - 12 minutes
- Rinse Cycle - 1 minute
- Rinse Cycle - 12 minute
- Rinse Cycle - 12 minutes
- Final Rinse - 2 minutes
- Rinse with Demineralized water 176°F (80°C) - 2 minutes
- Dy 1994°F (93°C) - 10 minutes

Dy 199.4°F (9°C) – 10 minutes
STERILIZATION RECOMMENDATIONS FOR REUSABLE INSTRUMENTS (GYNECARE TVT Introducer, GYNECARE TVT Rigid Catheter Guide)
The GYNECARETVT Introducer and GYNECARE TVT Rigid Catheter Guide are supplied non-sterile. To sterilize, steam autoclave prior to each use. Steam autoclave at a temperature of 27°F to 284°F (132°C to 140°C) for a minimum of 4 minutes (pre-vacuum). It is the responsibility of the end user to assure sterility of the product when using sterilization process recommended, since bioburden and sterilization equipment will vary.
INSTRIMENTA MAINTENAMCE

INSTRUMENT MAINTENANCE

GYNECARETYT Introducer

Before each use, inspect the threaded parts of the inner shaft.

GYNECARETYT Rigid Catheter Guide

Before each use, inspect the instrument. Check to ensure that the long end which traverses the catheter channel has no sharp edges or burrs.

HOW SUPPLIED

The GYNECARE TVT Device is provided sterile (ethylene oxide) for single use. Do not re-sterilize. Do not use if package is opened or damaged. Discard opened, unused devices. The reusable GYNECARE TVT Rigid Catheter Guide are supplied separately and are non-sterile. These accessories are to be cleaned and sterilized prior to each use as described above.

STORAGE

Recommended storage conditions for the GYNECARE TVT single use device are below 25°C, away from moisture and direct heat. Do not use after expiry date.

Caution: Federal (U.S.A.) law restricts this device to sale by or on the order of a physician.

Symbols Used on Labeling

2 = Do not reuse/resterilize

= Use by — year and month

STERILE EO = Method of Sterilization

CE-mark and identification number of Notified Body. Product conforms to the essential requirements of the Medical Device Directive 93/42/EEC

LOT = Batch number

See instructions for use